AMENDMENTS TO THE CLAIMS

1. (Currently amended) A process of preparing a compound [1-benzyl-4-[(5,6-dimethoxy-1-indanon)-2-yl]methylpiperidine] of the structural formula (II):

$$H_3CO$$
 CH_2
 $N-CH_2$

comprising catalytically hydrogenating a compound [1-benzyl-4-[(5,6-dimethoxy-1-indanon)-2-ylidene]methylpiperidine] of the structural formula (III):

$$H_3CO$$
 $N-CH_2$
 (III)

in the presence of a Raney nickel catalyst in a reaction solvent of <u>tetrahydrofuran</u>, toluene, <u>or a solvent mixture of toluene and an alcohol</u>, <u>in which appropriate quantities of soluble solvent is added</u>, <u>or tetrahydrofuran</u>, wherein the <u>reaction</u> solvent <u>is has a volume</u> 7 to 10 times of the volume of the compound of the structural formula (III).

2-6. (Canceled)

7. (Previously presented) The process according to claim 1, wherein the catalytic hydrogenation

is carried out at a hydrogen pressure of 0.05 to 7.0 MPa.

8. (Previously presented) The process according to claim 1, wherein the catalytic hydrogenation

is carried out at a hydrogen pressure of 0.1 to 1.5 MPa.

9. (Previously presented) The process according to claim 1, wherein the catalytic hydrogenation

is carried out at a hydrogen pressure of 0.5 to 1.5 MPa.

10. (Previously presented) The process according to claim 1, wherein a weight ratio of the

Raney nickel catalyst to the compound of the structural formula (III) is 3 to 30%.

11. (Previously presented) The process according to claim 1, wherein a weight ratio of the

Raney nickel catalyst to the compound of the structural formula (III) is 5 to 15%.

12. (Previously presented) The process according to claim 1, characterized in that the catalytic

hydrogenation is carried out at a reaction temperature of 4 to 60°C.

13. (Previously presented) The process according to claim 1, characterized in that the catalytic

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hydrogenation is carried out at a reaction temperature of about 4 to 40°C.

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14. (Previously presented) The process according to claim 1, characterized in that the catalytic hydrogenation is carried out at a reaction temperature of 10 to 25°C.

15. (**Currently amended**) A process for preparing a compound [1-benzyl-4-[(5,6-dimethoxy-1-indanon)-2-yl]methylpiperidine hydrochloride] of the structural formula (I):

$$H_3CO$$
 CH_2
 CH_2
 CH_2
 CH_2
 CH_2

comprising catalytically hydrogenating a compound [1-benzyl-4-[(5,6-dimethoxy-1-indanon)-2-ylidene]methylpiperidine] of the structural formula (III):

$$H_3CO$$
 $N-CH_2$
 (III)

in the presence of a Raney nickel catalyst in a reaction solvent of <u>tetrahydrofuran</u>, toluene, <u>or a solvent mixture of toluene and an alcohol</u>, in which appropriate quantities of soluble solvent is <u>added</u>, or <u>tetrahydrofuran</u>, wherein the <u>reaction</u> solvent is <u>has a volume</u> 7 to 10 times of the volume of the compound of the structural formula (III) to obtain a compound [1-benzyl-4-[(5,6-dimethoxy-1-indanon)-2-yl]methylpiperidine] of the structural formula (II):

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$$H_3CO$$
 CH_2
 $N-CH_2$

and then treating the compound of the structural formula (II) with hydrogen chloride or hydrochloric acid.

- 16. (New) The process according to claim 1, wherein the alcohol is methanol.
- 17. (New) The process according to claim 16, wherein the volume ratio of toluene to methanol in the solvent mixture is 4:1.
- 18. (New) The process according to claim 15, wherein the alcohol is methanol.
- 19. (New) The process according to claim 18, wherein the volume ratio of toluene to methanol in the solvent mixture is 4:1.